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Joseph Milanovich Loyola University Chicago, jmilanovich@luc.edu

William E. Peterman University of Missouri-Columbia

Kyle Barrett University of Georgia

Matthew Hopton United States Environmental Protection Agency

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Do species distribution models predict species richness in urban and natural green spaces? A case study using amphibians

Joseph R. MILANOVICH^{1*}, William E. PETERMAN^{2#}, Kyle BARRETT³⁺, and Matthew E. $HOPTON$ ^{1†}

¹U.S. Environmental Protection Agency, Office of Research and Development, National Risk Management Research Laboratory, Sustainable Technology Division, Sustainable Environments Branch, 26 West Martin Luther King Drive, Cincinnati, OH 45268, USA.

 2 Division of Biological Sciences, University of Missouri, Columbia, MO 65211

³D. B. Warnell School of Forestry and Natural Resources, University of Georgia, Athens, GA 30602, USA.

*Corresponding author: JRM, U.S. Environmental Protection Agency, Office of Research and Development, National Risk Management Research Laboratory, Sustainable Technology Division, Sustainable Environments Branch, 26 West Martin Luther King Drive, Cincinnati, OH 45268, USA, milanovich.joe@epa.gov, Phone: $011+1+(513)$ 569-7966.

- # WEP; bill.peterman@gmail.com
- + KB; kbarrett@warnell.uga.edu
- † MEH; hopton.matthew@epa.gov

Research Highlights:

- Amphibian species richness maps significantly over-predicted species richness. \overline{a}
- Over-prediction may have partially been a result of undersampling during surveys. $\frac{1}{2}$
- Over-prediction was likely due to poor model performance and undersampling. \blacksquare
- Despite over-prediction, models did project relative species richness well. \blacksquare

1 1. Introduction

2 Much of global population growth is occurring in urban areas (United Nations, 2004; Wu et al. 2011) where over one-half of the United States population resides in urban areas (MacKun $\&$ 3 Wilson, 2011). Nevertheless, human land use patterns are dynamic and some locations within 4 5 urban areas are experiencing declining populations. Such declines can result in land 6 abandonment and provide an opportunity to replace developed habitat with green infrastructure. It is well established that urbanization changes the biotic and abiotic properties of an ecosystem 7 and these impacts can reach far outside the urban area (Gaston, 2010). To reduce these effects, 8 9 there has been a movement to implement green infrastructure or incorporate green space to urban 10 areas. The benefits of green space in urban systems include increased psychological well-being, 11 recreational opportunities, and human health benefits (e.g., Barton & Pretty, 2010; Breuste & 12 Qureshi, 2011; Tzoulas et al., 2007; van den Berg, Hartig, & Staats, 2007). These benefits are often predicated by ecosystem services and functions such green space in urban ecosystems 13 provide (e.g., water filtration and quality; Bolund & Hunhammar, 1999; Faulkner, 2004; Gaston, 14 Davies, & Edmondson, 2010). Whereas urban habitats may not act as smaller versions of 15 16 undeveloped patches of land, they may still provide ecological and human-oriented benefits such 17 as providing habitat connectivity, which helps to sustain regional biodiversity (Goddard, Dougill & Benton, 2009; Irvine et al., 2010; Luck & Smallbone, 2010), or providing permeable surface 18 19 for stormwater infiltration, or water purification (Boyer & Polasky, 2004). Monitoring and 20 management for biodiversity has inherent value (Connery, 2009) and biodiversity conservation 21 within urban areas can help minimize extinction risk of some species and increase the value of 22 biota to humans as they more frequently encounter wildlife (Goddard et al., 2009). Toward this

23 end, metrics are needed to measure the degree to which urban green spaces sustain biota and 24 subsequent biodiversity.

An ideal metric would use taxa that serve as indicators of overall biodiversity, provide an 25 ecosystem service, and are a critical link to the biotic community within the green space (i.e., 26 27 provide ecosystem functions). Amphibians are often the most abundant, diverse group of 28 vertebrate organisms in forested and wetland systems, they serve as important food resources for 29 higher trophic levels, and in many systems are considered the top-predators (Burton $&$ Likens, 1975; Davic & Welsh, 2004; Gibbons et al., 2006). Amphibians are also considered to be 30 31 indicators of environmental stress (DeGarady & Halbrook, 2006; Southerland et al., 2004; Welsh 32 & Droege, 2001; Welsh & Ollivier, 1998), but see Kerby et al. (2010), and are known to provide 33 a number of ecosystem functions in natural ecosystems (Davic & Welsh, 2004; Regester, Lips & Whiles, 2006; Regester & Whiles, 2006; Whiles et al., 2006). Moreover, amphibians in urban 34 35 environments, like other biota, can enhance educational opportunities for human inhabitants 36 (Pickett et al., 2001). Because of their importance to ecosystems, ability to indicate environmental stress, and education value, research involving amphibians in urban systems is 37 38 warranted (Hamer & McDonnell, 2008; McDonnell & Hahs, 2008; Pickett et al., 2001; 39 Smallbone, Luck & Wassens, 2011). Due to time and financial constraints associated with conducting biotic surveys, modeling methods may provide assistance in understanding the value 40 41 of urban green space to this taxon.

42 Presence-only species distribution models (SDMs) are models that correlate species 43 distribution records to environmental data to predict areas of suitable habitat for taxa (see review in Guisan & Zimmerman, 2000; Guisan & Thuiller, 2005; Elith et al., 2006). They are a group 44 of approaches for identifying species distributions of undersampled species, predicting impacts 45

46 of environmental change on distributions, and identifying areas of conservation importance 47 (Elith & Leathwick, 2009). In recent years, validations of models in various forms have been increasing. For example, methods utilizing species occupancy or detection (Franklin et al., 2009; 48 Rota et al., 2011), independent and non-independent data validation (Araujo et al., 2005), and the 49 incorporation of field/survey data to inform or test model accuracy (Newbold et al., 2010; Pineda 50 51 & Lobo, 2009; Trotta-Moreu & Lobo, 2010) have been examined. However, studies simply using field data to validate whether models are projecting species distribution correctly are rare. 52 Others have noted several limitations to SDMs including exclusion of biotic, geographical, or 53 54 physiological constraints on species distributions, use of museum records that may be widely 55 variable in both spatial and temporal quality, and issues relating to extrapolation of model 56 predictions (see review in Elith & Leathwick, 2009). These limitations may be exacerbated 57 when modeling within spatial extents that include urban environments, because species are 58 sampled less in urban areas as ecologists tend to focus collections or research on natural areas (Gaston et al., 2010; Martin, Blossey & Ellis, 2012). Furthermore, lack of uniform sampling 59 across gradients of development presents a challenge to using SDMs in an urban landscape 60 61 because SDMs assume that biases in locality data (e.g., false absences) are not correlated with 62 environmental gradients used to build projected distributions (Bean, Stafford & Brashares, 2012; Hijmans, 2012). In addition, error in the predictions of SDMs varies over large spatial scales 63 (extent and resolution) due to increased spatial heterogeneity (Osborne, Foody & Suarez-Seoane, 64 2007; Smulders et al., 2010; Zhang & Zhang, 2007), such as variation of environmental, 65 66 landscape, and habitat structure. This trend may be seen at smaller spatial scales (extent and resolution) when using fine-scale data to build models (e.g., 30 m resolution) such as in urban 67 areas that have several classes of land use categories (e.g., habitat heterogeneity), as the 68

69 increased heterogeneity in urban areas within a smaller spatial scale could pose similar 70 prediction errors.

We tested whether species richness maps generated from SDMs can be used to prioritize areas 71 of high biodiversity value in urban and non-urban green space. We asked if SDMs built using 72 73 landscape variables associated with amphibian species richness could be used to project areas of 74 suitable habitat. We addressed this question by comparing modeled species richness maps (based on accumulated individual SDMs) to field surveys across a number of urban and non-75 urban green spaces. In addition, we investigated the landscape-level predictors of observed 76 77 amphibian species richness to determine what variables may be important to include or protect in 78 the creation, management, or conservation of urban and non-urban green space.

79

2. Materials and Methods 80

81 *2.1. Species Distribution Modeling using Maximum Entropy*

We developed species distribution models using Maxent version 3.3.3a (Phillips & Dudik, 82 2008) for 23 species of amphibians with current distributions within Hamilton County, Ohio, 83 84 U.S.A. Maxent is a software program that employs a machine learning method that is based on 85 the principle of maximum entropy to model species distributions using presence-only data coupled with environmental data. Entropy is characterized by Shannon (1948) as "a measure of 86 how much "choice" is involved in the selection of an event" and is utilized in the framework of 87 maximum entropy to examine species geographic distributions (Phillips, Anderson & Schapire, 88 89 2006). The approach estimates habitat suitability based on an input set of environmental 90 variables encompassing the region where a species is known to occur based on locality records. 91 The program maximizes the entropy in the probability distribution of suitability across all areas

92 of the distribution where empirical observations are lacking. For each species identified as 93 occurring in Hamilton County, OH, species presence data were obtained for the period of 1997– 2001from HerpNET (http://www.herpnet.org), Global Biodiversity Information Facility (Lane, 94 2003; GBIF; http://www.gbif.org), and personal collections of herpetologists (Appendix A). All 95 locality points were cross-referenced to each other and duplicate points were removed. 96 97 Furthermore, localities that fell outside the current species range (identified by county-level distribution maps found in Lannoo, 2005) were not utilized to develop models. To maximize 98 model quality, each model was built using at least 20 point locations for each species (Wisz et 99 100 $al., 2008.$ 101 We modeled the suitable habitat of each species across the National Hydrography Dataset 102 Plus (U.S. Geological Survey, 2005; NHDPlus; retrieved from http://www.horizon-103 systems.com/nhdplus/) Region 05 Unit B watershed delineation. This delineation was necessary 104 to encompass the environmental variability of each species and provide predicted records from across a larger geographic region adjacent to our primary study area. Additionally, amphibian 105 distributions can be restricted by geographic barriers (i.e., large rivers), and this region 106 107 encompassed a large portion of each species range, as well as the primary area of our field 108 surveys used to validate our models. This watershed delineation also encompasses a number of similar adjacent urban areas (e.g., Lexington, KY, Indianapolis, IN, and Huntington, WV; 109 110 Appendix B) that could be examined in future studies. 111 We chose 11 initial environmental variables that others have shown to be important to 112 amphibians (e.g., Herrmann et al., 2005; Weyrauch & Grubb Jr, 2004) and had available data. 113 Geographic data layers were at a resolution of 30 m and variables extracted from these included land cover, elevation, canopy cover, distance from stream, and slope/aspect. We created three 114

115 layers using a moving window analysis to compute a ratio of cells classified as urban or water 116 within moving windows of 150, 300, and 500 m. Lastly, we calculated the number of cells representing wetland habitat, as defined by the National Wetlands Inventory layer (U.S. Fish and 117 Wildlife Service, 2011; retrieved from http://www.fws.gov/wetlands/), within 300 m and 2 km 118 119 moving windows. We tested collinearity of the 11 layers by extracting environmental 120 information from 1000 randomly selected points (selected using the Random Point Generation 121 Tool within the Hawth"s Analysis Tool software for ArcGIS) within the watershed. A 122 correlation matrix was generated and correlations with $r \ge 0.70$ were considered highly 123 correlated. When pairs of variables exceeded this threshold $(r \ge 0.70)$, we chose one variable 124 from the pair that we considered the most biologically relevant. Eight variables (listed below 125 with the data source) were chosen for inclusion in the final distribution model. Land cover 126 (Homer et al., 2007; retrieved from http://www.mrlc.gov/) was reclassified into one of seven 127 classes as follows (original land cover classes given in parenthesis): "forest" (41, 42, and 43), water/wetlands (11, 90, and 95), , low intensity developed" (21 and 22), , medium intensity 128 developed" (23), "high intensity developed" (24), "natural non-forest" (31, 52, and 71), and 129 130 "agriculture" (81 and 82). Canopy cover values were taken from the National Land Cover 131 Database (NLCD; Homer et al., 2007; retrieved from http://www.mrlc.gov/). Elevation data were derived from NHDPlus (National Hydrography Dataset; U.S. Geological Survey, 2005; 132 133 retrieved from www.horizon-systems.com/nhdplus/). The distance from stream (DSL) was 134 measured as distance (in meters) from the nearest drainage. Drainages were delineated from the 135 flow accumulation layer (downloaded from NHDPlus; retrieved from www.horizon-136 systems.com/nhdplus/) as areas that drained 100 ha or more. The synthetic slope/aspect layer 137 (TASL; which represents aspect ranging from -1 [NE] to 1 [SW] and is weighted based on the

161 a continuous probability of suitable habitat for each species, it is logistically unfeasible to present 162 each location as a probability of occupancy; therefore, we converted the continuous suitability surface $[0-1$ from Maxent to presence/absence $(1/0)$] using four model output thresholds applied 163 by Maxent; fixed cumulative value 10 (FC10), minimum training presence (MTP), 10 percentile 164 training presence (10% TP), and maximum training sensitivity plus specificity (MTSPS). Next, 165 166 we generated four species richness maps based on the accumulated binary modeled distributions of each species using the four Maxent thresholds. This four-threshold approach makes our 167 results comparable to other studies that provide predictions based on strict environmental 168 169 distributions of species (*i.e.*, thresholds that maximize the agreement between observed and 170 predicted distributions; Cramer, 2003). This approach allows us to present model predictions 171 that relax the assumption of strict environmental control on species" distributions, and provides a 172 range of scenarios that could influence the predicted suitable habitat at the species-level and 173 comprehensive species richness.

We used null models to test the significance of each species distribution model (Raes $&$ ter 174 Steege, 2007). We generated five null data sets, each with 1000 sets of sample points that were 175 176 randomly drawn without replacement from the pool of 620 background points. We generated a 177 null data set with the number of random points per distribution equal to 20, 45, 75, 150 and 250 data points, which represented a range of presence points available to model each species. 178 179 Maxent was used to calculate the area under the curve (AUC) for the 1000 null data sets to create 180 an AUC frequency distribution. The calculated AUC for each species model was compared to the 95th percentile AUC value of the null frequency distribution created from the representative 181 182 number of sample points (20, 45, 75, 150, or 250). A species model performs better than random 183 (e.g., null model) and is considered significant if the calculated AUC is greater than the

corresponding 95th percentile AUC of the null-distribution (Raes & ter Steege, 2007). 184

185 *2.2. Examining differences in amphibian species richness*

186 *2.2.1. Amphibian surveys*

To evaluate the degree to which SDMs accurately predict amphibian richness, we conducted 187 188 amphibian surveys at 20 sites across the Cincinnati, OH metropolitan area following methods 189 proposed in Shaffer et al. (1994). Each site consisted of identified green space, spanned a range of sizes $(3-1758 \text{ ha})$. Appendix C), and were spread across a gradient of urbanization (e.g., within 190 191 and outside the designated Cincinnati, OH metropolitan area; Fig. 1; Appendix B). Surveys were 192 conducted March – June 2011 and each site was sampled three times with a 3–5 week period 193 between repeat samples. To standardize our search effort within each site, nine plots were 194 selected and surveyed at each site. We attempted to place three plots in each of three habitat 195 types associated with amphibian richness: Wetlands/ponds, streams, and terrestrial (forest or non-forest natural [e.g., grassland/prairie]) habitat. In the event that three each of 196 wetlands/ponds or streams were not available, we increased the number of terrestrial plots to 197 198 reach nine total plots for each site. If more than three of an aquatic habitat type were available 199 (e.g., wetlands/ponds or streams) we sampled plots with the most suitable amphibian habitat (i.e., forested wetlands/ponds or streams). We employed three survey techniques at each site: area 200 201 sensitive dip-net surveys for pond and/or ephemeral wetland habitats, area sensitive terrestrial 202 (searching cover objects) surveys, and leaf litter bags and dip-netting area sensitive surveys for 203 headwater streams following Chalmers & Droege (2002) and Waldron et al. (2003). At each wetland/pond, we dip-netted several 1 m² areas adjacent to the bank using a 40.6 X 40.6 cm (3.2) 204 205 mm mesh size) net. Plots were placed every 10 m until the entire perimeter of the body of water

was sampled. For terrestrial plots, we surveyed five, 5 m² sub-plots at each site. Plots were 206 chosen by placing a single 5 $m²$ plot within the center of each habitat type occurring on the site 207 (forest or non-forest natural [prairie or grassland habitats]) and an additional 5 m^2 plot in each 208 cardinal direction 25 m from the center plot. Within each sub-plot, all cover objects and existing 209 leaf litter were searched. Within each stream, five 1 m^2 plots were created every 10 m starting 210 211 approximately 20 m from the confluence of a larger stream, road, or trail and continuing upstream. Within each plot, we placed one leaf litter bag $(55 \times 25 \text{ cm} \text{ made with } 1.3 \text{ cm}^2 \text{ mesh})$ 212 filled with deciduous leaf litter from the surrounding forest in a wetted portion of the stream 213 214 channel with a large rock on top to prevent dislodging. We followed the methods of Nowakowski & Maerz (2009) and Peterman, Truslow, & Samuel (2008) to remove contents 215 from each bag; we thoroughly dip-netted the 1 $m²$ plot by dislodging substrate parallel to the 216 217 stream bank across the width of the stream. In addition to our structured surveys, any animals 218 encountered (seen or heard) while travelling between habitat types or plots within a site were noted. All animals captured or heard (frog calls) were identified to species. 219

220 *2.2.2. Validation of amphibian surveys and richness comparisons*

221 We examined how well predicted species richness matched observed species richness by 222 comparing the maximum species richness value from each site between observed and predicted (from each Maxent threshold) values derived from our maps. We compared differences between 223 species richness within "urban" and "non-urban" sites by categorizing a site as "urban" if $\geq 50\%$ 224 225 of the adjacent landscape within a 2-km buffer of the centroid of the site had urban cells (NLCD 226 categories $21 - 24$) based on our NLCD 2001 reclassified maps. To test for differences between 227 predicted species richness from each model threshold, we used paired *t*-tests to compare 228 observed and predicted (from each threshold) species richness values from each site surveyed.

229 To examine if we adequately sampled species richness, we estimated species richness using 230 EstimateS version 8.2 (Colwell, 2009) for six sites that had been surveyed on at least 10 occasions. Additional survey data were derived from reports summarized in Davis, Krusling, 231 and Ferner (1998). These reports summarized data from county-park-level surveys that took 232 place between 1988 and 1995, surveyed a similar level of habitat heterogeneity and utilized 233 234 similar survey methods to our study, and lasted approximately one year. We chose to make comparisons between species richness at this group of sites using the Chao 2 estimator because 235 236 this algorithm produces species accumulation curves that approach maximum values with few 237 samples (i.e., the estimator was the least sensitive to undersampling; Colwell & Coddington, 238 1994). The Chao 2 algorithm inflates the observed species richness by a factor derived from the 239 number of species observed only once or twice within a total sample. The estimator is calculated as $S_{Chao2} = S_{obs} + Q_1^2 / 2Q_2$, where S_{obs} is the observed species richness and Q_1 and Q_2 are the 240 241 number of species detected only once or twice per site, respectively. Thus, this estimator accounts for the fact that species differ in detectability and uses the relative frequencies of 242 species that are rarely detected to estimate the number of taxa that are present but not detected 243 244 (Chao, 1987; Colwell & Coddington, 1994). For each additional site we calculated coverage 245 (number of species observed/number of species estimated X 100), exclusive species (percentage of species only observed in a given site or category), and completeness of a sample (species 246 observed as a percentage of the total number of species expected in the site; Gardner et al., 247 248 2007). This approach estimated the amphibian species richness at sites visited ≥ 10 times and 249 provided an estimate of the number of sampling periods needed to accurately capture the 250 estimated richness value. These values were then compared to the observed amphibian species

251 richness values derived from the current surveys and the surveys of Davis, Krusling, and Ferner 252 $(1998).$

253 *Identifying actors that influence observed species richness*

We used a general linear model and an information theoretic approach to examine which 254 environmental factors were most predictive of observed amphibian species richness. This model 255 256 is specific to existing green spaces, unlike the species distribution models, which provide a more general estimate of species richness patterns. As a result, the environmental variables used in 257 this analysis are similar to those in the distribution model; however, some differences do exist. 258 259 Observed species richness was the dependent variable. Independent variables included the 260 percent of urban land within a 2-km buffer of the centroid of each site (square root transformed), 261 the number of wetlands within a 2-km buffer of the centroid of each site (log transformed), 262 percent forest within each site (square root transformed), and percent non-forest natural habitat 263 (agriculture and non-forested natural habitat) within each site (square root transformed) as continuous predictor variables. Using the same variables as above, we evaluated the importance 264 of landscape variables on observed species richness using Akaike Information Criterion 265 266 (Burnham & Anderson, 2002). These variables were derived from our reclassified NLCD 2001 267 land-cover layer and a National Wetlands Inventory layer (U.S. Fish and Wildlife Service, 2011; http://www.fws.gov/wetlands/) and were chosen out of an initial set of 11 independent variables 268 that we thought would have an influence on amphibian species richness (Table 3). We generated 269 270 a correlation matrix of all 11 variables and chose four variables that were not correlated with 271 each other ($r \le 0.70$). Variables were transformed to meet normality assumptions (Sokal & 272 Rohlf, 1995).

273 Lastly, to determine the relationship between our observed species richness values and the 274 mean predicted species richness values across the urban gradient, we conducted an analysis of covariance. Maxent predictions (predicted species richness) and field observations (observed 275 species richness) were factors in the analysis, percent urbanization within 2-km of each site 276 277 (square root transformed) was our covariate, and observed species richness and mean predicted 278 species richness values were used as our dependent variables. The significance of the interaction 279 term was evaluated to determine if the relationship between observed and predicted species richness estimates differed across the urban treatments. 280

281

282 3. Results

283 The mean AUC for amphibian distribution models based on landscape variables was 0.83 $(\text{range} = 0.70 - 0.96; \text{ median} = 0.83; \text{ Appendix A})$ and the AUC for each species was better than 284 random (i.e., model AUC values exceeded the 95th percentile of the null AUC distributions). 285 Observed and predicted species richness varied across each site (Appendix C). Mean observed 286 species richness was triple in non-urban sites compared to the value in urban sites (mean species 287 288 richness values; overall = 3.2, urban = 1.9, non-urban = 6.2). There were significant differences 289 between observed and predicted species richness for each threshold (Table 1; all *t*-tests $P \leq$ 0.001). Overall, mean differences between observed and predicted species richness within each 290 291 threshold ranged from 253 to 558% (mean range $= 8$ to 18 species). These differences increased 292 when "non-urban" sites were isolated from "urban" sites (mean range $=$ 343 to 619%; mean 293 range $= 8$ to 17 species) and decreased when sites sampled greater than 10 times were isolated 294 from sites sampled three times (mean range = 14 to 98%; mean range = 2 to 11 species; see 295 "Additional" column in Table 1).

296 The MTSPS threshold, our most conservative threshold, had the lowest levels of over-

297 prediction between observed and predicted species richness in both urban and non-urban sites; whereas the MTP threshold had the greatest levels of over-prediction. The species accumulation 298 curves (Fig. 2) for all green spaces sampled > 3 times, except Farbach-Werner Nature Preserve, 299 suggest these sites had been (more or less) exhaustively sampled (i.e., the species accumulation 300 301 curve reaches a plateau). There was no site for which the final estimate of species richness from 302 the species accumulation curves was achieved by the third sample (based on, for example, Chao 2 richness estimator; Fig. 2; Table 2). 303

304 Observed amphibian species richness varied depending on the number of wetlands and 305 percent of urbanization within a 2-km buffer around the centroid of each site (Tables 3 and 4). 306 Percent of forest or non-forest natural habitat within each site did not predict observed species 307 richness (Table 3). The best model (based on lowest AICc and highest w_i) for observed species 308 richness also includes number of wetlands and percent of urbanization within a 2-km buffer around the centroid of each site (Table 4). Percent of forest or non-forest natural habitat within 309 each site did not predict observed species richness (Table 3). Despite the significant 310 311 discrepancies in absolute richness values between the observed and predicted species richness 312 values, there was a significant trend of decreasing richness with increasing urban habitat $(P \leq$ 0.001 for both observed and mean predicted richness). Further, the slopes of these two trends 313 314 did not differ $(P = 0.913$; Appendix D) between modeled and observed values and the 315 relationships between observed and predicted species richness was significantly correlated to the 316 level of adjacent urbanization (Appendix D).

317

4. Discussion and Conclusions 318

319 The use of SDMs built with high-resolution landscape variables and real-time amphibian 320 surveys allows us to assess whether this technique is a valid method to identify areas of higher and lower species richness across urban and non-urban green space. Our approach of modeling 321 followed by field surveys revealed differences between observed and predicted species richness 322 across urban and non-urban green space (Table 1). These differences can be interpreted as (1) 323 324 sites were undersampled, thus our observed measure of species richness could be inaccurate, or (2) the SDMs are over representing actual species distributions – which accumulate to unrealistic 325 predictions of species richness across the landscape. Our results suggest the differences are 326 327 likely due to both factors. First, species accumulation curves derived from the Chao 2 estimator 328 based on six sites that were extensively surveyed (i.e., ≥ 10 sampling events over approximately 329 one year), indicated three sampling periods was not a sufficient effort to capture species richness. 330 In each of these six sites, species accumulation curves did not plateau until between six to 10 331 sampling periods (Fig. 2). We conducted an intensive single-season survey that reasonably 332 mimics the effort many agencies would implement to understand factors driving species distributions of amphibians. We know from studies with a more intensive sampling effort that 333 334 this is not sufficient to capture all species, because often species are unavailable for capture for 335 entire breeding years (e.g., Barrett & Guyer, 2008).

However, undersampling alone did not explain all of the differences between observed and 336 predicted species richness. Chao 2 species richness estimates showed SDMs still over-predicted 337 338 species richness even in sites considered to be adequately sampled (Table 2). For that subset of 339 sites, we have reasonable confidence (empirically demonstrated via richness estimator) that all 340 differences between sampled and predicted species richness are not due to undersampling alone. This suggests the differences are at least partially a consequence of SDM predictions. 341

342 We propose the SDM influence on over-prediction is the result of the inherent nature of 343 SDMs. By solely using environmental variables to construct predictions of a species" suitable habitat, SDMs fail to incorporate biological or geographic influence on species distributions 344 (e.g., Guisan & Thuiller, 2005; Heikkinen et al., 2006; Luoto et al., 2005), also referred to as 345 realized versus fundamental niche (Rodda, Jarnevich & Reed, 2011; Soberón & Peterson, 2005). 346 347 This in turn can lead to an overestimation of species suitable habitat, as only areas of suitable habitat, not realized distribution, are projected. This overestimation may increase when adding 348 349 multiple individual models to create species richness maps (Hortal & Lobo, 2006; Pineda & 350 Lobo, 2009; Thullen, Sartoris & Walton, 2002; Vasconcelos, Rodriguez & Hawkins, 2011), as 351 was the case in this study. Furthermore, although the use of target-group background data is 352 suggested to reduce sampling bias (Phillips et al., 2009), use of this approach could be a possible 353 source of overestimation. Models developed utilizing replicate samples and random background 354 data could result in a greater or lesser over-estimation of species richness values. Further investigation into the influence of various modeling approaches to overestimation of species 355 richness is warranted. 356

357 Using multiple thresholds to develop a range of scenario's of species predicted suitable 358 habitat was a definite strength of this study (Fig. 1; Appendices B and C), and the differences between predicted species richness for different model thresholds should not be ignored. The 359 360 importance of presenting a range of model thresholds has been supported in other studies (Liu et 361 al., 2005; Milanovich et al., 2010; Pineda & Lobo, 2009; Thullen et al., 2002). We suggest 362 presenting a range of thresholds allowed the geographic range of suitable habitat for each species 363 to be both larger and smaller than realized ranges. This may account for some of the geographic and biotic influences on species ranges, however, as our results indicate, threshold adjustment 364

365 may be a necessary, but not entirely sufficient means for generating more precise SDMs. 366 Despite the over-prediction of our models and differences between model thresholds, we argue this technique is a useful tool for management of green space by identifying areas suitable 367 for amphibians in urban ecosystems. Specifically, although the model richness estimate was 368 high, there was no difference between the trend in observed richness and modeled richness 369 370 across the urban treatments. Species distribution models are increasingly used to identify potential areas for conservation and management of biodiversity (Lawler, Wiersma & 371 Huettmann, 2011; Loiselle et al., 2003). In this study, SDMs identified areas of higher 372 373 suitability, as green space with higher observed species richness generally had higher predicted 374 species richness (Appendix B and C). For example, the Cincinnati Nature Center and 375 Richardson Nature Preserve (the non-urban sites with the highest observed species richness) and 376 Farbach-Werner Nature Preserve and Triple Creek Park (the urban sites with the highest 377 observed species richness) each had some of the highest predicted species richness within each 378 threshold (Appendix C). One commonality between these sites is each site had a high number of adjacent wetlands (within 2-km of the centroid), and therefore, a high number of aquatic habitats 379 380 surveyed (Appendix C). With respect to management of existing or development of new urban 381 green spaces, we found the number of wetlands adjacent to each site was a significant predictor of observed amphibian richness (Tables 3 and 4). Our predictive models support the importance 382 383 of wetlands in urban green space to amphibian species richness, as the two urban sites with 384 surveyed wetlands (Farbach-Werner N.P. and Triple Creek Park) had the highest predicted 385 species richness of all urban sites. This highlights the importance of prioritizing the 386 maintenance, restoration, or mitigation of wetlands in urban green space. Furthermore, this is a 387 testament to the usefulness of SDMs for identifying areas of potential conservation importance,

388 but emphasizes that models must be constructed with biologically relevant variables that 389 facilitate presence of a particular species or taxon (e.g., wetlands or forest for amphibians). In conclusion, we tested whether species distribution models could be utilized to identify 390 areas of higher amphibian species richness in urban ecosystems. If one"s goal is to make rapid 391 392 decisions concerning the management or creation of green space or infrastructure based on 393 conserving areas of higher biodiversity, this method provides relative measures of richness. Our results indicate substantial effort is needed to sufficiently survey and identify areas of suitable 394 habitat for amphibians in urban and non-urban green spaces. Field surveys often require 395 396 expertise, time, energy, and funds; whereas SDMs can be developed using existing, readily 397 available data (e.g., museum records) and can be conducted using a variety of freely available 398 programs (Elith et al., 2006). We offer two suggestions on how to improve SDM use if one"s 399 goal is to identify areas important to high biodiversity. First, the development of species-specific 400 models, particularly in cases where there may be a desire to increase the abundance of a particular species of concern, could enhance predictive ability. These models could utilize key 401 (spatially explicit) variables important to the biology of a particular species. Second, we 402 403 recommend using SDM approaches that incorporate an estimation of probability of occurrence 404 and species detectability, where such data are available for species of interest. This method has been successfully utilized and detailed in more recent studies (e.g., Franklin et al., 2009; 405 406 Newbold et al., 2010; Rota et al., 2011). Unfortunately, our data did not permit such analyses. 407 We suggest incorporating one or both of these approaches to strengthen the accuracy of SDMs in 408 predicting species richness.

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Table 1. Estimates of how much SDMs over-predicted species richness values in green spaces in and around Hamilton County, Ohio. Values in the table represent the percentage each threshold $(MTP = \text{minimum training presence threshold}, MTSPS = \text{maximum training sensitivity plus}$ specificity threshold, 10% TP = 10 percentile training presence threshold, FC10 = fixed cumulative value 10 threshold) over-predicted species richness compared to surveyed values for all sites (overall), urban sites (sites with $\geq 50\%$ urbanization within the 2-km buffer of centroid), non-urban sites (sites with $\leq 50\%$ urbanization within 2-km buffer of centroid), and from eight sites where past species richness values (derived from county-park surveys) were added to our current species richness values derived from current surveys (labeled Additional).

Table 2. Summary species richness data for amphibians at sites sampled ≥ 10 times in the Hamilton County, Ohio area.

Table 3. Results from a general linear model investigating the factors that influenced the observed amphibian species richness within each site and associated parameter estimates and 95% confidence intervals (CI). Percentage data were square root transformed and number of wetlands within 2-km buffer was log transformed. Excluded correlated variables not chosen for analysis were as follows: size; standard deviation of land cover within parks and 2-km buffer of centroid; percent forest, non-forest natural, and urban cells within 2-km buffer of centroid, and number of aquatic habitats surveyed within each site.

Table 4. Best Akaike Information Criteria (AIC) models for observed species richness across all 20 green spaces. Shown are only the confidence set of candidate models (i.e., models with AIC w_i within 10% of highest model). Variables included were percent urbanization within 2-km buffer, number of wetlands within 2-km buffer, percent forest within site, and percent non-forest natural habitat within each site.

Table 1. Estimates of how much SDMs over-predicted species richness values in green spaces in and around Hamilton County, Ohio. Values in the table represent the percentage each threshold $(MTP = \text{minimum training presence threshold}, MTSPS = \text{maximum training sensitivity plus}$ specificity threshold, 10% TP = 10 percentile training presence threshold, FC10 = fixed cumulative value 10 threshold) over-predicted species richness compared to surveyed values for all sites (overall), urban sites (sites with $\geq 50\%$ urbanization within the 2-km buffer of centroid), non-urban sites (sites with \leq 50% urbanization within 2-km buffer of centroid), and from eight sites where past species richness values (derived from county-park surveys) were added to our current species richness values derived from current surveys (labeled Additional).

Table 3. Results from a general linear model investigating the factors that influenced the observed amphibian species richness within number of wetlands within 2-km buffer was log transformed. Excluded correlated variables not chosen for analysis were as follows: each site and associated parameter estimates and 95% confidence intervals (CI). Percentage data were square root transformed and Size; standard deviation of land cover within parks and 2-km buffer of centroid; percent forest, non-forest natural, and urban cells within 2-km buffer of centroid, and number of aquatic habitats surveyed within each site.

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Figure 1. (a) Distribution of locality points (black dots) used to model species suitable habitat within the watershed boundary. The boundary is located across several states with Ohio and Kentucky (the states where the surveys were conducted) located in the northeast and southern border of the map; and (b) the three counties where each green space (site) was surveyed; metropolitan areas are outlined with a red line and surveyed green spaces are represented by filled green polygons. Surveys were conducted throughout various locations within each site. The four lower panels represent the predicted amphibian species richness maps within the three counties surveyed as derived from the maximum training sensitivity plus specificity threshold (MTSPS) (c), minimum training presence threshold (MTP) (d), 10 percentile training presence threshold $(10\% \text{ TP})$ (e), and fixed cumulative value 10 threshold (FC10) (f)

Figure 2. Sample-based species accumulation curves for the six heavily sampled sites in the Hamilton County, Ohio metropolitan area

Appendix A. Area under curve (AUC) values.

- Appendix B. Species richness maps.
- Appendix C. Summary data.
- Appendix D. Results from analysis of covariance.

Appendix A. Area under curve (AUC) values for each species modeled.

Appendix B. Depictions of: a) Cities of Indianapolis, Indiana (1), Cincinnati, Ohio (2) Lexington, Kentucky (3), and Huntington, West Virginia (4) located within the watershed boundary. Predicted amphibian species richness maps within the watershed boundary derived from the maximum training sensitivity plus specificity threshold (b), minimum training presence threshold (c), 10 percentile training presence threshold (d), and fixed cumulative value 10 threshold (e).

Appendix D. Results of ANCOVA and regression analyses to determine if species richness observations/estimates differed across the urban gradient. Species richness was the dependent variable, which was estimated either through field-observations (observed SR) or Maxentpredictions (predicted SR). Square root transformed percent urbanization within 2 km of site was the covariate in the analysis. The non-significant interaction term indicates that the relationship between species richness estimates and urbanization was not different. Estimates for each method differed significantly, and urbanization had a significant, negative effect on richness $r^2 = 0.627, P$ $r^2 = 0.491, P$ Predicted species richness = $23.765 - 0.925*%$ urbanization.

Figure 1.

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